Cell Lines Used for Research Only

HCT 116 (3D)

Cat. No. : CL-0096-3D

Origin and General Characteristics

Cell Name	HCT 116 (3D)
Organism	Human
Tissue	Colon
Morphology	Tight 3D spheres
Growth Properties	Suspension spheres
Background	HCT 116 (3D) tumorspheres are cancer stem cell-enriched spheroids obtained from HCT 116 cells by enrichment and culture with specialized medium.
Biosafety Level	BSL-1

Culture Conditions and Handling

Complete Medium	Complete culture medium for HCT 116 (3D)[CM-0096-3D]
Subcultivation Ratio	0.5×10^5 cells/mL
Medium Renewal	2 to 3 times per week
Dissociation Duration	3-5 min
Incubation Atmosphere	Air, 95%; CO_2, 5%
Temperature	37°C
Freeze medium	Freezing Medium (Serum-free & animal origin-free)[PB180438]
Storage Conditions	For long-term cryopreservation, cryovials should be stored in liquid nitrogen at −150°C to −196°C. Storage at −80°C is restricted to short-term interim use only.
Instructions	1. Check all containers for leakage or breakage. 2. Remove the frozen cells from the dry ice packaging and immediately transfer them to liquid nitrogen (liquid or vapor phase) for long-term cryopreservation. 3. When cultured in non-low-adherence vessels, tumor spheres may grow with partial surface attachment.
Subculturing Procedure	1. Transfer the tumor spheres to a 15 mL centrifuge tube, centrifuge, and discard the supernatant. 2. Add approximately 1 mL of Accutase Cell Detachment Solution (Cat. No.: PB180201), and gently invert the tube to ensure all tumor spheres are fully submerged. 3. Incubate the tube at 37°C for 3–5 min, gently pipette the suspension up and down several times, and verify under a microscope that the tumor spheres have dissociated into a single-cell suspension. 4. Collect the cell suspension, centrifuge at 1200 rpm for 3 min, then aspirate and discard the supernatant. 5. Add HCT 116 (3D Spheroid-Specific) Cell Complete Medium, gently pipette to achieve a homogeneous cell suspension, seed the cells at a density of 50,000 cells/mL into a new culture flask, and bring the

mixture to the final volume with fresh medium. Incubate with either a loosened cap or a vented cap.

Handling Recommendations for Cryopreserved Cells

Preparations Before Thawing

1. Prepare the recommended complete culture medium and preheat it to 37°C for 30 minutes.
2. Prepare 9 mL of complete medium within a sterile centrifuge tube to dilute the cryoprotectant present in the frozen cell suspension.
3. Use sterile gloves to protect the cryovial from contact with the water bath water, and preventing cell contamination.

Thawing Procedure

1. Retrieve the cryovial containing frozen cells from liquid nitrogen storage. Place it in sterile gloves and immediately submerge it in a 37°C water bath.
2. Thaw the cells rapidly (<1 minute) by gently swirling the vial in the 37°C water bath until only a small ice crystal remains.
3. Transfer the vial to a biosafety cabinet. Wipe the exterior of the vial with 75% ethanol before opening.
4. Transfer the thawed cells dropwise into a preheated 2 mL centrifuge tube containing complete medium.
5. Centrifuge the cell suspension at approximately 250 × g for 3 minutes.
6. Aspirate and discard the supernatant, then resuspend the cells in 1 mL of HCT 116 (3D) complete medium.
7. Seed cells at a density of 50,000 cells/mL into 125 mL shake flasks (or low-adherence culture plates / low-adherence T25 flasks), add 10–20 mL of HCT 116 (3D) complete medium, mix thoroughly, and incubate at 37°C with 5% CO_2.

Notes

1. The entire recovery process should be completed as quickly as possible.
2. Select the culture vessel size based on the number of cells in the cryovial. Procell single cryovials are recommended for resuspension in T25 flasks or 10 cm dishes.
3. Minimize the time thawed cells are kept at room temperature. Cryoprotectant must be immediately diluted or removed by centrifugation.

Precautions After Receipt of Frozen Cells

1. Upon unpacking, inspect the condition of the frozen cells and dry ice, and take photographs immediately. The following after-sales service will be provided based on these photographs, including assessment of the remaining dry ice, verification that the cryovial was fully buried in dry ice, and evaluation of whether the cells thawed and refrozen during transit.
2. Upon receipt, the cells should be transferred to liquid nitrogen immediately or directly resuscitated. If liquid nitrogen is unavailable, the cells can be temporarily stored at -80°C, however, the storage period should be limited to less than two weeks whenever possible. Prolonged storage at -80°C may gradually reduce post-thaw cell viability, and the extent of this viability loss is unpredictable.
3. Ensure that the operator has sufficient knowledge and experience in cell culture, and the laboratory is equipped with essential instruments, including a biosafety cabinet, CO_2 incubator,

inverted microscope, centrifuge, water bath. Carefully review the cell instruction manual to understand key cell characteristics including growth properties (adherent or suspension), morphology, basal medium requirements, serum concentration, cytokine supplementation, subcultivation ratio, medium renewal schedule.

4. After resuscitation, observe the cells under a microscope and record the cell status by taking photographs (1-3 images each at 100× and 200× magnification for 3 consecutive days). These images will serve as supporting documentation for follow-up services. In addition, a small aliquot of cells may be used to assess cell viability by automated cell counting or trypan blue staining.
   Note: Cells should not be observed too frequently within the first 24 hours after resuscitation, as this may affect cell growth or adherence. Observation once per day is sufficient.
5. After sufficient expansion, the cells should be cryopreserved as soon as possible to ensure availability of backup stocks for future experiments or in case of unexpected experimental failure. Cells in the logarithmic growth phase should be selected for cryopreservation. After cryopreservation, it is recommended to resuscitate one vial to verify post-thaw viability and confirm successful cryopreservation.